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## **Committee on Technical Barriers to Trade**

Original: English

## **NOTIFICATION**

## Addendum

The following communication, dated , is being circulated at the request of the delegation of <u>Brazil</u>.

Title: Draft Resolution number 760, December 27th, 2019.

Reason for Addendum:	
[]	Comment period changed - date:
[]	Notified measure adopted - date:
[]	Notified measure published - date:
[]	Notified measure enters into force - date:
[]	Text of final measure available from <sup>1</sup> :
[]	Notified measure withdrawn or revoked - date: Relevant symbol if measure re-notified:
[X]	Content or scope of notified measure changed and text available from¹: <a href="https://antigo.anvisa.gov.br/documents/10181/5740831/RDC_931_2024pdf/42c43a_0e-99cd-4114-9fe1-d5b6421a68d1">https://antigo.anvisa.gov.br/documents/10181/5740831/RDC_931_2024pdf/42c43a_0e-99cd-4114-9fe1-d5b6421a68d1</a> New deadline for comments (if applicable):
[]	Interpretive guidance issued and text available from <sup>1</sup> :
[]	Other:

**Description:** Resolution 742, 10 August 2022 - previously notified through G/TBT/N/BRA/955/Add.1 - which establishes the minimal technical requirements for relative bioavailability and bioequivalence studies that supports dossier of consent for clinical research, market

authorization or post-market authorization of medicines, in the terms of this resolution, was change by Resolution 931, 09 October 2024.

The final text is available only in Portuguese and can be downloaded at:

https://antigo.anvisa.gov.br/documents/10181/5740831/RDC 931 2024 .pdf/42c43a0e-99cd-4114-9fe1-d5b6421a68d1

<sup>1</sup> This information can be provided by including a website address, a pdf attachment, or other information on where the text of the final/modified measure and/or interpretive guidance can be obtained.